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## Amendments to the Claims:

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Please amend the claims as follows:

1-60. (cancelled)

A lipid liposomal formulation containing a compound that 61. (currently amended)

is:

a diester of a compound of formula A (i)

YCO-NHCHCO-G\*

CH<sub>2</sub>-Z-X

where:

each ester is 1-25C;

YCO is  $\gamma$ -glu or  $\beta$ -asp;

G\* is phenylglycine;

Z is CH2, O or S; and

X is a hydrocarbon radical which is alkyl (6-8C), benzyl, or naphthyl; or a pharmaceutically acceptable salt thereof; or

a compound of formula I (ii)

where:

R<sub>1</sub> and R<sub>2</sub> are each independently linear or branched alkyl (1-25C), cycloalkyl (6-25C), heterocycle (6-25C), ether or polyether (3-25C), or R<sub>1</sub> and R<sub>2</sub> together have 2-20 C atoms and form a macrocycle with the remainder of formula I; and

X is as defined above for formula A;

or a pharmaceutically acceptable salt thereof;

where the lipids of the lipid liposomal formulation are consist of egg phosphatidylcholine

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and egg phosphatidylglycerol in a ratio of 0.75-1.25:0.75-1.25 by weight and the ratio of lipids to compound is 3.5-4.5:0.5-1.5 by weight.

- 62. (currently amended) The <u>lipid liposomal</u> formulation of claim 61 where the compound is γ-glutamyl-S(benzyl)cysteinyl-R-phenylglycine diethyl ester or a pharmaceutically acceptable salt thereof.
- 63. (cancelled)
- 64. (currently amended) The <u>lipid liposomal</u> formulation of claim 63 61 where the ratio of lipids to compound is 3:1-6:1 by weight.
- 65. (canceled)
- 66. (currently amended) The lipid liposomal formulation of claim 61, having
- (i) at least 50% degree of encapsulation of the compound; and
- (ii) an average vesicle size of 50-2000 nm.
- 67. (currently amended) The <u>lipid liposomal</u> formulation of claim 66 where the degree of encapsulation is above 80%.
- 68. (currently amended) The <u>lipid liposomal</u> formulation of claim 66 where the vesicle size is 400-600 nm.
- 69. (currently amended) The <u>lipid liposomal</u> formulation of claim 61 which is a <u>liposomal formulation composed consists essentially</u> of 1 part compound, 2 parts egg phosphatidylcholine, 2 parts eg
- 70. (currently amended) The lipid liposomal formulation of claim 69 which comprises liposomes composed consists essentially of 1 part  $\gamma$ -glutamyl-S(benzyl)cysteinyl-R-phenylglycine diethyl ester or a pharmaceutically acceptable salt thereof, 2 parts egg phosphatidylcholine, 2 parts egg phosphatidylglycerol, and 7 parts sucrose by weight.

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The lipid liposomal formulation of claim 70 which 71. (currently amended) comprises lyophilized liposomes composed consisting essentially of 1 part  $\gamma$ -glutamyl-S(benzyl)cysteinyl-R-phenylglycine diethyl ester or a pharmaceutically acceptable salt thereof, 2 parts egg phosphatidylcholine, 2 parts egg phosphatidylglycerol, and 7 parts sucrose by weight.

A method of preparing a lipid liposomal formulation 72. (currently amended) containing a compound that is:

a diester of a compound of formula A (i)

where:

each ester is 1-25C;

YCO is  $\gamma$ -glu or  $\beta$ -asp;

G\* is phenylglycine;

Z is  $CH_2$ , O or S; and

X is a hydrocarbon radical which is alkyl (6-8C), benzyl, or naphthyl; or a pharmaceutically acceptable salt thereof; or

a compound of formula I (ii)

where:

R<sub>1</sub> and R<sub>2</sub> are each independently linear or branched alkyl (1-25C), cycloalkyl (6-25C), heterocycle (6-25C), ether or polyether (3-25C), or R<sub>1</sub> and R<sub>2</sub> together have 2-20 C atoms and form a macrocycle with the remainder of formula I; and

X is as defined above for formula A;

or a pharmaceutically acceptable salt thereof;

which method comprises formulating the compound in a lipid liposomal composition

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where the lipids of the lipid liposomal formulation are consist of egg phosphatidylcholine and egg phosphatidylglycerol in a ratio of 0.75-1.25:0.75-1.25 by weight and the ratio of lipids to compound is 3.5-4.5:0.5-1.5 by weight.

The method of claim 72 where the compound is  $\gamma$ -73. (previously presented) glutamyl-S(benzyl)cysteinyl-R-phenylglycine diethyl ester or a pharmaceutically acceptable salt thereof.

74. (cancelled)

The method of claim 72, further comprising extrusion. 75. (previously presented)

The method of claim 72, further comprising lyophilization. 76. (previously presented)

The method of claim 72 which comprises dissolving 1 part 77. (previously presented) 7-glutamyl-S(benzyl)cysteinyl-R-phenylglycine diethyl ester or a pharmaceutically acceptable salt thereof, 2 parts egg phosphatidylcholine, and 2 parts egg phosphatidylglycerol in ethanol/water, injecting the solution into water containing 7 parts sucrose, and extruding to form a liposomal formulation.

The method of claim 77 which comprises dissolving 1 part 78. (previously presented) Y-glutamyl-S(benzyl)cysteinyl-R-phenylglycine diethyl ester or a pharmaceutically acceptable salt thereof, 2 parts egg phosphatidylcholine, and 2 parts egg phosphatidylglycerol in ethanol/water, injecting the solution into water containing 7 parts sucrose, extruding to form a liposomal formulation, and lyophilizing the liposomal formulation to form lyophilized liposomes.

A lipid liposomal formulation prepared by the method of 79. (currently amended) claim 77.

80. (currently amended) A lipid lyophilized liposomal formulation prepared by the method of claim 78.

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81. (currently amended) A method for modulating hematopoiesis or protecting against the destructive effects of chemotherapy comprising administering to a subject in need thereof a lipid liposomal formulation according to any one of claims 61, 62, 64, 66 to 71, 79, and 80.